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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁵ : C07K 15/06, 15/14, 15/12, C12P 21/08, G01N 33/577, A61K 27/02, 39/395	A1	(11) International Publication Number: WO 94/21686 (43) International Publication Date: 29 September 1994 (29.09.94)
(21) International Application Number: PCT/AU94/00139 (22) International Filing Date: 21 March 1994 (21.03.94) (30) Priority Data: PL 7926 19 March 1993 (19.03.93) AU (71) Applicant (for all designated States except US): NORTHERN SYDNEY AREA HEALTH SERVICE [AU/AU]; Level 3 - Vindin House, Royal North Shore Hospital, Pacific Highway, St Leonards, NSW 2065 (AU). (71)(72) Applicant and Inventor: SINOSICH, Michael, Joseph [AU/AU]; 9 Sirius Place, Engadine, NSW 2233 (AU). (74) Agent: GRIFFITH HACK & CO.; P.O. Box 4164, Sydney, NSW 2001 (AU).		(81) Designated States: AT, AU, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, ES, FI, GB, GE, HU, JP, KP, KR, KZ, LK, LU, LV, MG, MN, MW, NL, NO, NZ, PL, PT, RO, RU, SD, SE, SI, SK, TT, UA, US, UZ, VN, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i>
(54) Title: PAPP-A, ITS IMMUNODETECTION AND USES (57) Abstract This invention concerns PAPP-A, its immunodetection and the clinical benefits of such immunodetection. Specifically, the invention includes monoclonal antibodies against PAPP-A and the use of these antibodies to detect PAPP-A at a very early stage of pregnancy. The invention also covers the use of the monoclonal antibodies for the detection of specific types of cancer and Down's Syndrome pregnancies.		

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PAPP-A, ITS IMMUNODETECTION AND USESTechnical Field

The present invention relates to: purified pregnancy-associated plasma protein-A (PAPP-A); PAPP-A variants; polynucleotides encoding PAPP-A; isolation and purification of PAPP-A; monoclonal antibodies raised against PAPP-A; use of these monoclonal antibodies and PAPP-A for diagnostic purposes, including a kit for the assaying of PAPP-A levels in a sample; use of PAPP-A measurement, in conjunction with another marker of trophoblastic activity (such as chorionic gonadotrophin or its subunits), to discriminate between feto-placental abnormalities, such as Trisomy 21 (Down Syndrome), and oncological status, such as Gestational Trophoblastic Diseases (GTD); use of PAPP-A monoclonal antibodies to isolate fetal trophoblast cells for prenatal fetal cytogenetic diagnosis; use of PAPP-A, as a target antigen for active immunological contraception, and PAPP-A antibodies as a passive contraceptive vaccine; use of PAPP-A as a medicament and a medicament comprising an effective amount of PAPP-A.

BACKGROUND ART

Placental proteins are those proteins expressed during pregnancy by the human placenta. The ability to detect the presence and concentrations of these proteins has the potential to provide a reliable diagnostic marker of fertilisation, implantation and pregnancy prognosis.

A number of placental proteins have now been isolated and at least partially characterised. These include - human chorionic gonadotropin (hCG), pregnancy-specific β_1 - glycoprotein (SP1), placental protein 5 (PP5), early pregnancy factor (EPF), and pregnancy-associated plasma protein-A (PAPP-A)¹.

These proteins are detectable, in maternal blood, at various stages during pregnancy. For example, EPF activity is detectable within 24 hours after conception. HCG is measurable just after implantation, at about 9 to 11 days post-ovulation, SP1 is detectable from 18 to 23

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days post-ovulation. In singleton pregnancies, PAPP-A can be detected approximately 28-32 days post-ovulation².

Placental proteins are also detectable for varying periods during pregnancy. For example, EPF is detectable at least for the first half of pregnancy, whereafter activity declines until it is totally absent during the third trimester in some women. HCG levels rise rapidly to peak at about 8 to 12 weeks gestation. The levels of SP1 rise exponentially with peak concentrations being reached at term pregnancy. Like SP1, PAPP-A concentrations also rise exponentially in the first trimester of pregnancy to peak at term².

Whilst it has been suggested to measure the presence of placental proteins for early detection of pregnancy (for example, see European Application 316919), there is a growing body of documented evidence that at least some placental proteins, particularly PAPP-A, may be used to predict pregnancy viability, including early pregnancy failure, extra-uterine gestations, aneuploid and/or abnormal pregnancies, such as Down's Syndrome³ and Cornelia de Lange Syndrome⁴.

PAPP-A, first described almost two decades ago¹, is a large zinc containing glycoprotein, rich in carbohydrate, with many physicochemical similarities to a 2-macroglobulin⁵. It has been detected in maternal circulation⁵, pre-ovulatory ovarian follicular fluid⁶, in seminal plasma⁷ and blood of patients with trophoblastic disease⁸.

PAPP-A is a homotetramer, with each monomeric subunit having a molecular weight of approximately 200 kDa. The subunits are linked by disulphide bonds to form dimers of approximately 400 kDa. Native PAPP-A consists of two dimers linked by Van der Waals (ionic) forces. Native PAPP-A has a molecular weight of approximately 820 kDa, regardless of whether it is derived from follicular fluid, seminal plasma, oncological or normal placental tissue⁷. The mature protein has a 2- β 1 electrophoretic mobility, with an isoelectric point of approximately

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4.2-4.5⁹. It is a non-competitive and potent inhibitor of human granulocyte elastase¹⁰.

It has been suggested that the biological function of PAPP-A is to act as a local protective barrier against host (maternal) phagocytic-proteolytic defences to either insemminated sperm or the developing feto-placental unit¹¹. This may be due to PAPP-A forming a protective sheet around the chorionic villus at the utero-placental interface¹². Disruption of this protective layer may explain the correlation between depressed PAPP-A levels and pregnancy failure. PAPP-A may also play a role in zinc homeostasis⁵.

Schindler and Bischof¹³ suggested that the protein was ubiquitous and, therefore, of little practical use in pregnancy viability diagnosis.^{14,15} However, it was subsequently shown that these results were due to impure PAPP-A isolates, due to the difficulties in isolating PAPP-A free of a 2-macroglobulin, and polyspecific antisera¹⁶.

Sinosich et al.³ first suggested that a depressed or undetectable PAPP-A level in maternal blood was diagnostic of pregnancy failure. Later, Sinosich et al.¹⁷ showed that, of five successful in vitro fertilisation volunteers, three patients with normal pregnancy outcome had circulating PAPP-A levels within the 80% confidence limits of the normal range. By contrast, circulating PAPP-A levels in the patient who spontaneously aborted at seventeen weeks were below the tenth percentile throughout the entire gestation. In the fifth patient, who had a ruptured ectopic pregnancy, PAPP-A could not be detected at any stage during the pregnancy. These findings were complimented by Westergaard et al. (1983)¹⁸, who reported that, in a sample group of 51 patients, who conceived spontaneously, with vaginal bleeding in the first half of pregnancy, concentrations of PAPP-A were consistently lower in pregnancies which failed. Similarly, Sinosich et al. (1985)¹⁹ showed that, in a group of 21 women who conceived by in vitro

fertilisation, PAPP-A levels were consistently depressed, for many weeks, in those women whose pregnancies failed. The same group showed that, of forty seven serum samples obtained from patients with a tubal pregnancy, only two
5 were positive for PAPP-A, indicating that severely depressed or undetectable serum PAPP-A levels were an aid in the diagnosis of extra-uterine pregnancy.²⁰

In 1990, Brambati *et al.*²¹ reported that first trimester maternal serum concentrations of PAPP-A were
10 low in pregnancies associated with Down's syndrome. Later, Wald *et al.* (1992)³ confirmed that PAPP-A concentration was significantly lower in women with Down's syndrome pregnancies compared to PAPP-A levels in a control group of normal pregnancies.

15 It has also been reported that PAPP-A was detected in the circulation of patients with hydatiform mole², suggesting a potential role for PAPP-A quantification in diagnosis and management of certain tumours.

These findings demonstrate the potential diagnostic
20 value of measuring PAPP-A levels for monitoring fetoplacental status. Moreover, in view of the increasing use of *in vitro* fertilisation techniques, and the relatively high proportion of early pregnancy failures associated with these techniques, the measurement of
25 PAPP-A levels to monitor pregnancy viability and thereby minimise patient trauma is clinically advantageous.

Lin. *et al.* (1974)²² described the use of an electroimmunoassay to measure PAPP-A levels in advanced pregnancy. This assay was insensitive and limited to
30 latter stages of pregnancy. Sinosich *et al.* (1982)² and (1984)⁶ described the first sensitive radioimmunoassay (RIA) which detected PAPP-A in serum obtained from first trimester pregnancies. This assay used radioactively
35 labelled purified PAPP-A together with rabbit anti-human PAPP-A antiserum. The sensitivity of this RIA (2.9 µg/L) enabled PAPP-A detection in maternal blood after the first six weeks of pregnancy. The assay made it possible to detect PAPP-A in other fluids (amniotic fluid, seminal

plasma, follicular fluid, gestational trophoblastic disease, culture media), previously beyond the limits of detection. This assay also made it possible to study the kinetics and physiology of maternal PAPP-A levels in the first trimester of pregnancy, a crucial stage for fetoplacental development.

The PAPP-A RIA differed from standard protocols in that molecular size of tracer (^{125}I -PAPP-A; Mr 820kDa) and immune complexes required modification of the separation phase. Optimal separation of antibody-bound from antibody-free tracer was achieved with second antibody - 7.5% (w/v) polyethylene glycol (PEG) solution, in the ratio of 2:1, 2nd antibody-PEG to assay reaction volume. Under these conditions, assay blank values could be reduced to 5-7%, whilst maximum binding would approach 60-70%.

The development of sensitive and reliable techniques for measuring PAPP-A is dependent on being able to isolate the protein in a sufficiently pure form and/or the generation of monospecific antibodies.

A number of methods for isolating and purifying PAPP-A have been described previously. For example, Lin. et al. (1974)²³ describe a procedure based on classical protein fractionation technology. This procedure utilised:

- i) solubility;
- ii) charge; and
- iii) size.

This procedure was technically cumbersome and resulted in a low yield of impure material.

Bischof (1979)²⁴ described a method of isolating and purifying PAPP-A. This procedure utilised:

- i) solubility;
- ii) charge;
- iii) lectin affinity;
- iv) size fractionation; and
- v) negative immunoaffinity.

This was even more cumbersome than the earlier procedure. The final product was still impure and yields

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remained low.

Sutcliffe et al. (1979)²⁵ and Folkersen et al. (1981)²⁶ described a purification method which used;

- i) solubility;
- 5 ii) positive immunoaffinity chromatography (immobilised anti-PAPP-A antibodies); and
- iii) size or charge, respectively.

Step (ii) is detrimental to PAPP-A integrity and neither procedure resulted in pure PAPP-A. Although the final
10 yield was improved by this method, the quality of the protein yield was sacrificed.

Based on an interaction with heparin (Sinosich et al. 1981)²⁷, Sinosich et al. (1982)¹⁰ reported the first application of heparin-Sepharose for PAPP-A purification.
15 For the first time it was possible to prepare a high yield (22%) of highly purified PAPP-A. However, by more current and stringent criteria, this preparation was also found to be impure.

Disclosure of the Invention

20 The inventor has developed a purification procedure which enables PAPP-A to be purified to a much higher degree than achieved with prior art purification techniques and in particular to a level which permits the amino-acid sequencing of the protein. The inventor has
25 sequenced the first thirteen (13) N-terminal amino acids of human PAPP-A and this sequence is:

Glu-Ala-Arg-Gly-Ala-Pro-Glu-Glu-Pro-Ser-Pro-Pro-Ser

According to a first aspect of the present invention there is provided the protein PAPP-A, substantially free
30 of other (primate) proteins, said PAPP-A being a homotetramer having a molecular weight of approximately 820 kiloDaltons and an isoelectric point of approximately 4.5, each monomeric subunit of the homotetramer having a molecular weight of approximately 200 kiloDaltons and an
35 N-terminal amino-acid sequence of Glu-Ala-Arg-Gly-Ala-Pro-Glu-Glu-Pro-Ser-Pro-Pro-Ser.

The ability to sequence the protein indicates that the preparation is at least 90% pure.

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Biologically active proteins are usually only present in trace amounts in biological systems. Consequently, purification from naturally occurring sources is expensive and time consuming. Having purified
5 PAPP-A to sufficient purity to obtain amino acid sequence for the protein, it is now possible for the first time to produce degenerate nucleotide probes which correspond to the determined amino acid sequence. These probes can then be used to detect nucleic acid encoding PAPP-A and
10 clone it. At this level of purity PAPP-A can also be used as an antigen to raise antibodies to detect PAPP-A expressing clones.

Production of the PAPP-A protein by recombinant techniques and isolation of the DNA encoding PAPP-A also
15 makes it possible to obtain a complete amino acid sequence for PAPP-A. The complete amino acid sequence can be used to determine appropriate sites for mutagenesis in the production of clinically useful PAPP-A variants.

20 According to a second aspect of the present invention there is provided a PAPP-A variant. Variants of PAPP-A in accordance with this invention are polypeptides which correspond to or comprise a portion of PAPP-A or have homology with the PAPP-A amino acid
25 sequence.

For the purposes of this description "homology" between two peptide sequences connotes a likeness short of identity, indicative of a derivation of the first sequence from the second. In particular, a polypeptide
30 is "homologous" to PAPP-A of the invention if a comparison of amino-acid sequences between the polypeptide and PAPP-A reveals an identity of greater than about 70%. Such a sequence comparison can be performed via known algorithms, which are readily
35 implemented by computer.

According to a third aspect of the invention there is provided a polynucleotide encoding PAPP-A or a PAPP-A variant of the present invention.

According to a fourth aspect of the invention, there is provided a method for isolating and purifying PAPP-A, said method comprising:

- (i) applying the sample to a positive affinity chromatography on heparin-Sepharose;
- (ii) size fractionating the fractions obtained from step (i) on a gel filtration column; and
- (iii) applying the fractions obtained from step (ii) to an anion exchange column; and
- (iv) applying the fractions obtained from step (iii) to a matrix with immobilized antibodies.

According to a fifth aspect of the invention, there are provided monoclonal antibodies which recognise PAPP-A. Typically the monoclonal antibodies of this aspect recognise PAPP-A but do not significantly cross-react with immobilized heparin-Sepharose binding proteins extracted from normal male serum. Preferably the monoclonal antibodies are specific for PAPP-A. In particular, the inventor has developed five (5) monoclonal antibodies which specifically recognise five (5) distinct epitopes of the PAPP-A protein.

The first monoclonal antibody - clone 25-1 - has been isotyped as IgG₂ (K light chain) with an affinity constant of 1.26×10^{-9} . Clone 25-1 reacts specifically with human PAPP-A but not with chimpanzee or rhesus PAPP-A.

The second monoclonal antibody - Clone 27-66, isotyped as IgM with K light chain specifically recognises PAPP-A from human, chimpanzee and rhesus primates.

The third monoclonal antibody - Clone 5-62, isotyped as IgG₁ with K light chains with an affinity constant of 2.65×10^{-9} reacts specifically with human and chimpanzee PAPP-A but not with rhesus PAPP-A.

The fourth monoclonal antibody - Clone 18-9, isotyped as IgG with K light chains, reacts specifically with human and chimpanzee PAPP-A but not with rhesus

PAPP-A.

The fifth monoclonal antibody - Clone 41.1, has yet to be isotyped and characterised.

The advantages conferred by the development of monoclonal antibodies to PAPP-A are that they are specific for a distinct epitope. This specificity allows the development of an assay with increased sensitivity and reliability. This increased sensitivity permits the detection of PAPP-A in the circulation within the first six weeks of normal pregnancy and depressed levels in abnormal pregnancies.

The use of monoclonal antibodies also permits the development of immunoassays which are independent of labelled antigen (PAPP-A). The term "immunoassays", as used herein, includes any method for PAPP-A detection mediated by the use of antibodies (classes IgG, IgA, IgD, IgE or IgM) or derivatives thereof (including Fab and F(ab)₂ fragments). Antibodies may be polyclonal or monoclonal, and may be generated by active immunisation of a host animal (including mouse, rat, guinea pig, rabbit, sheep, horse, donkey or other mammalian species), hybridoma technology and/or molecular biology for production of recombinant antibodies. Quantitative and qualitative PAPP-A detection may be achieved in liquid and/or solid phase and, in addition to immunoassays, includes in situ hybridisation and immunohistochemistry, by way of example.

Monoclonal antibodies also allow the development of sandwich assays with greater amplification potential. The single epitope specificity of the antibodies permits the detection of PAPP-A active sites and, therefore, quantification of bio-active PAPP-A.

According to a sixth aspect of the present invention, there is provided a sample method for the detection of PAPP-A in a sample using a monoclonal antibody of the fifth aspect to detect PAPP-A. Typically, the method comprises the steps of:-

(i) contacting the sample with a PAPP-A capture

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phase which consists of one or more of the following:

- (a) monoclonal PAPP-A antibody;
- (b) polyclonal PAPP-A antibody;
- (c) immobilised heparin;
- 5 (d) immobilised divalent metal cations (Cu^{++} , Zn^{++} , Co^{++} , Ni^{++});
- (e) immobilized lectins (Concanavalin A, Helix pomatia, Lens culinaris, Limulus polyphemus, phytohaemagglutinin, Ricinus communis, Wheat
10 germ, or others with equivalent specificities), and,
- (f) Other specific (such as receptors) or non-specific ligands (such as dyes) with which PAPP-A has an affinity.
- 15 (ii) contacting the immobilised or captured PAPP-A sample with a monoclonal antibody, of the fifth aspect of the invention, labelled with a detectable marker;
- (iii) incubating the sample and the labelled antibody to permit the labelled antibody to bind to any
20 PAPP-A in the sample and;
- (iv) detecting the labelled antibody.

Typical methods include enzyme immunoassays (EIAs) or immunoradiometric assay (IRMA) formats.

According to a seventh aspect of the present
25 invention there is provided a kit for the detection of PAPP-A in a sample which kit comprises:

at least one monoclonal antibody according to the fifth aspect of the present invention together with a positive and/or negative control. The PAPP-A assay kit,
30 developed for manual and/or automated application, is not limited to human application. Selection of appropriate polyclonal-monoclonal PAPP-A antibody combination can extend application to sub-human primates.

The ability to produce large quantities of
35 recombinant PAPP-A or variants thereof permits use as a medicament for the treatment of pregnant women with low or zero PAPP-A levels in order to increase the likelihood of the pregnancy proceeding successfully to term.

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According to an eighth aspect of the present invention, there is provided the use of PAPP-A as a medicament together with a pharmaceutically acceptable excipient to treat a pregnant patient exhibiting the
5 absence of or low levels of PAPP-A.

Formulation of PAPP-A for this purpose with standard carriers, excipients and diluents is performed in accordance with standard pharmaceutical techniques.

According to a ninth aspect of the present invention
10 there is provided a medicament comprising an effective amount of PAPP-A together with a pharmaceutically acceptable carrier, diluent or excipient.

According to tenth aspect of the present invention, there is provided the use of PAPP-A as a target for
15 immunological contraception. Immunological fertility regulation may be achieved by active vaccination against intact PAPP-A or fragments thereof. Active vaccination may be achieved by any of the accepted oral, mucosal or subcutaneous routes (of antigen administration) used to
20 induce an immune response in humans. The PAPP-A antigen may be chemically modified, genetically engineered or the genome inserted into a vector to enhance immunogenicity when expressed in the host.

Administration of PAPP-A antibodies (or fragments
25 thereof) may be applied for fertility regulation by passive immunisation or for immunolocalisation and/or immunoneutralisation of trophoblastic tumors or management of extrauterine pregnancies. Antibody administration can be achieved by any of the procedures
30 outlined for PAPP-A antigen administration.

According to the eleventh aspect of the present invention, there is provided the use of PAPP-A (polyclonal and monoclonal) antibodies for isolation of trophoblast cells from systemic maternal circulation or
35 reproductive tract (vaginal cavity, cervical canal, cervical os). The isolated trophoblast cells may be cultured and applied for prenatal fetal karyotyping.

According to the twelfth aspect of the present

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invention, there is provided the use of PAPP-A immunodetection to discriminate between normal and abnormal pregnancies. Abnormal pregnancies includes multiple gestations (more than one fetus), extra-uterine implantation, anembryonic pregnancies, death in utero, incomplete miscarriage, spontaneous abortion, fetal malformations and aneuploidies, as examples. Quantitative or qualitative PAPP-A immunodetection may be performed on maternal biological fluids, such as blood. Blood may be collected by routine venepuncture, finger pricking or by any accepted medical procedure. The blood may be anticoagulated or coagulated, to permit the removal of supernatant for PAPP-A analysis. The blood may be stored frozen or dried onto an inert absorptive medium, such as filter paper, for transportation, subsequent extraction and analysis.

According to the thirteenth aspect of the present invention, there is provided the use of PAPP-A immunodetection in female biological fluids, such as ovarian follicular fluid and reproductive tract secretions, to assess folliculogenesis, granulosa cell status and ovulation.

According to the fourteenth aspect of the present invention, there is provided the use of PAPP-A immunodetection for diagnosis and management of gestational trophoblastic disease (GTD). In hydatidiform mole, a benign GTD exhibits a molar expression ratio of PAPP-A to hCG (or free b-subunit) which is indistinguishable from that observed in normal pregnancy. As the GTD transforms into the more aggressive and invasive choriocarcinoma, the molar expression ratio of PAPP-A to hCG (or free b-subunit) decreases, that is hCG (or free b-subunit) expression is maintained or increased, whereas, PAPP-A expression is downregulated. This trophoblast antigen expression ratio can be applied as an algorithm for diagnosis and management of GTD.

According to a fifteenth aspect of the invention, PAPP-A antibodies are used for the immuno-treatment of

trophoblastic tumors.

According to a sixteenth aspect of the present invention, there is provided the use of PAPP-A immunodetection in male reproductive tract secretions, such as seminal plasma and prostatic fluid, to assess the clinical state of male accessory glands, such as prostate gland, of the reproductive tract.

According to a seventeenth aspect of the present invention, there is provided the use of PAPP-A immunodetection for specific discrimination of pregnancies carrying Trisomy 21 (Down Syndrome) fetuses from normal and abnormal pregnancies. Although both hCG (and free b-subunit) and PAPP-A are trophoblast proteins, in this particular clinical situation their respective expression rates are discordant. Whereas PAPP-A expression is downregulated, hCG (and free b-subunit) expression is upregulated. Therefore, the molar expression of PAPP-A to hCG (or free b-subunit) can be applied as an algorithm for prenatal screening of Down Syndrome pregnancies in early gestation.

Brief Description of the Drawings

The invention will now be described with reference to the following drawings:

- Figure 1 - The N-terminal amino acid sequence of PAPP-A.
- Figure 2 - Anion exchange on Mono Q. Purification of primate (human, A; macaque, B) PAPP-A by high pressure liquid chromatography anion exchange on Mono Q. Dashed lines represent increases in NaCl concentration to effect PAPP-A desorption (at 46-48 min).
- Figure 3 - Assay protocol for PAPP-A quantification consists of a capture phase immobilized on an inert support and a monoclonal antibody for quantification. The monoclonal antibody may be tagged or visualized with a secondary antibody. The latter option

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enhances signal amplification and assay sensitivity.

- Figure 4 - Influence of capture antibody concentration on PAPP-A standard curves.
- 5 Figure 5 - Influence of temperature on the binding equilibrium of immobilised immunoglobulin at 4°C, room temperature and 37°C, Maximal PAPP-A capture was achieved after overnight incubation (15 hours).
- 10 Figure 6 - Binding equilibrium of monoclonal antibody clone 25-1 at various concentrations. Binding equilibrium was achieved by 4 hours at room temperature.
- Figure 7 - Standard curves for PAPP-A dose response at two dilutions for clone 25-1.
- 15 Figure 8 - Logit-log graphs for human serum and normal and caesarian placental homogenates assayed by RIA with clone 25-1. The results suggest parallelism between circulating and tissue PAPP-A.
- 20 Figure 9 - Logit-log by graphs for human serum and normal and caesarean placental homogenates assayed by EIA with clone 25-1. The results again suggest parallelism between circulating and tissue PAPP-A.
- 25 Figure 10 - Logit-log graphs for human, chimpanzee and rhesus placental homogenates assayed by EIA with clone 27-66. The results indicate that clone 27-66 equally reacts with PAPP-A from all three species.
- 30 Figure 11 - Logit-log graphs for human, chimpanzee and rhesus placental homogenates assayed by EIA with clone 5-62. The results indicate that clone 5-62 reacts with PAPP-A from human and chimpanzee but not rhesus.
- 35 Figure 12 - Logit-log graphs for human, chimpanzee and rhesus placental homogenates assayed by EIA with clone 25-1. The results indicate

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that clone 25-1 reacts only with human PAPP-A.

- Table 1 - Monoclonal 5-62 and 25-1 were tested for competition against a variety of antisera.
- 5 Table 2 - The results indicate that the monoclonals recognise different epitopes.

Best Mode for Carrying out the Invention

Methods of recombinant PAPP-A production and preparation of variants are performed in accordance with standard techniques as taught, for instance, in Maniatis et al. (1984)³¹ which is herein incorporated by reference. Formulation of PAPP-A medicaments is performed in accordance with standard pharmaceutical techniques.

Where recombinant techniques are used for the preparation of the protein of the invention, the protein can be prepared by constructing degenerate DNA probes coding for amino acid sequence of PAPP-A as described above and using these probes to isolate positive clones from c-DNA libraries according to standard techniques as found in text books such as Maniatis et al. (1984)³¹. These positive clones are then incorporated into appropriate vectors with expression cassettes according to standard techniques. These vectors are then used to transform host cells to allow the production of the recombinant PAPP-A protein.

The selection of appropriate vectors, hosts and expression strategies can be performed in accordance with standard techniques of molecular biology.

Suitable expression systems include, for instance, Chinese hamster ovary expression systems.

It will be readily apparent to a skilled addressee that where PAPP-A is produced recombinantly it may not be necessary to purify the protein before use. Recombinant production can be conducted in well-defined prokaryotic and eukaryotic hosts. Production in non-human systems yields the protein in a form at least substantially free of other human proteins.

The homologous polypeptides can be produced by

conventional site-directed mutagenesis, which is one avenue for routinely identifying residues of the molecule that can be modified without rendering the resulting polypeptide biologically inactive, or by chemical
5 synthesis.

Those variants which correspond to or comprise a portion of PAPP-A of the invention without being coincident with PAPP-A of the invention, within the scope of the invention, are those molecules which retain the
10 immunogenic or biological activity of the native PAPP-A protein.

These variants may be prepared synthetically by peptide synthesis techniques, recombinantly or by cleavage from an isolated protein of the invention.

15 Methods of isolating, characterising and cloning polynucleotides encoding PAPP-A using probes constructed from the N-terminal amino acid sequence are performed in accordance with standard techniques as taught, for instance, in Maniatis et al (1982)³¹.

20 Isolation and Purification of PAPP-A

The protocol consists of five chromatographic procedures and is applicable to human and non-human PAPP-A species. The protocol can be used to isolate PAPP-A from any biological fluid, culture medium or tissue
25 extract.

1. Positive Affinity Chromatography on Heparin-Sepharose:

The column was equilibrated with aqueous buffers, such as 50mM Tris-HCl, pH 7.4-7.8, containing 150 mM NaCl (TBS - 0.15M NaCl). Where was sample taken from ? Sample
30 application (1ml/min) resulted in only 11.3% of applied serum proteins being bound to the matrix. The remaining proteins do not interact with heparin and are easily removed from PAPP-A. By increasing the ionic strength of equilibration buffer to BS - 0.3M NaCl, the low affinity
35 heparin interaction was reduced and only 0.5% of applied proteins were retained by the ligand (heparin). Although PAPP-A recovery was 100%, the yield was 39% for a

purification factor of 486.

Heparin-PAPP-A interaction is heterogeneous, with minor amounts of PAPP-A not interacting with heparin (Sinosich 1985)²⁷. Although the significance of this heterogeneity is uncertain, it is clearly not related to heparin. Therefore, the heterogeneity resides with PAPP-A and may be related to PAPP-A isoforms or metabolic clearance, and, hence, provide a distinction between functional and inactive PAPP-A.

Desorption of matrix bound PAPP-A was achieved with application of high ionic strength buffers, such as TBS containing 0.6M NaCl or greater (up to 2.0M NaCl). PAPP-A elutes as a sharp peak and is concentrated into 3 or 4 fractions, dependent on column size and fraction volume. Desorption may also be achieved by applying the heparin antagonist, protamine sulphate, but matrix regeneration is more complex.

2. Size Fractionation:

PAPP-A containing fractions were pooled (36ml) and applied onto a 5 x 90cm column (Vol=1.71l) packed with gel filtration matrix Ultragel Aca34 or Aca22 or equivalent. PAPP-A containing fractions were pooled and dialysed against 20 mM TBS, containing 10 mM NaCl, pH 7.4-7.8. PAPP-A recovery for this procedure was 98.8% with an overall purification factor of 827.

3. Anion Exchange (Mono Q):

The dialysed PAPP-A pool (110 ml) from size fractionation was applied onto a 10 ml anion exchange column, such as Mono Q, by high pressure liquid chromatography. The PAPP-A containing pool from size fractionation was applied in 10 ml aliquots at 1ml/min. After 20 min., buffer ionic strength was increased to TBS-0.5M NaCl by a gradual linearly increasing concentration gradient in NaCl. This removed low affinity interactions. The high affinity PAPP-A interaction was dissociated with a stepwise increase in NaCl concentration to TBS-1M NaCl (Figure 2).

4. Negative Immunoaffinity Chromatography:

Rabbit antibodies against serum proteins, heparin binding serum and tissue proteins were immobilised onto inert particles such as CNBr-Sepharose. PAPP-A containing fractions (25 ml) from step 3 (above) were repeatedly passed over this matrix to remove any contaminants.

5. Positive Affinitive Chromatography:

This is optional, but provides an easy means of concentrating PAPP-A from larger into smaller volumes prior to aliquotting and storage. PAPP-A pool (90 ml), from step 4 was applied onto a 120 ml Heparin-Sepharose column. Matrix bound PAPP-A is desorbed with TBS-1M NaCl for a final PAPP-A yield of up to 22%, for an overall purification factor of 1483.

6. The purity of PAPP-A preparation was assessed and confirmed by:

- i) SDS-PAGE analysis (Sinosich et al., 1990)²⁸
- ii) radioimmunoassay (Sinosich and Zakher, 1991²⁹;
Sinosich et al. 1982², 1984⁶) and
- iii) NH₂-terminal amino acid sequencing.

The NH₂-terminal amino acid sequence for PAPP-A was sequenced by the School of Biological Sciences, Macquarie University according to standard techniques and was determined to be:

Glu-Ala-Arg-Gly-Ala-Pro-Glu-Glu-Pro-Ser-Pro-Pro-Ser-

As there is only one NH₂-terminal amino acid sequence detected, this indicates both that PAPP-A preparation is free of contamination with other proteins, and PAPP-A subunits are identical.

As this sequence is unique to PAPP-A, immunodetection of PAPP-A means the detection of a protein or proteins containing this NH₂ terminal amino acid sequence.

35 Monoclonal Production

Immunisation:

Pure PAPP-A, isolated from pooled late pregnancy serum was used to immunise BALB/c mice by two protocols.

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PAPP-A (30 μ g) in 0.05M sodium phosphate buffer, pH 7.4, containing 150 mM NaCl (50 μ l; PBS) was emulsified in 50 μ l Complete Freund's Adjuvant (CFA) and injected into the footpads of mice. Fourteen (14) days later, a
5 booster dose of 65 μ g PAPP-A/100 μ l PBS was injected into the thigh muscle of Mouse 1. This mouse was sacrificed 5 days later for hybridoma cell fusion.

Mouse 2 was subcutaneously immunised with 130 μ g PAPP-A in CFA, followed, three weeks later, with an
10 intraperitoneal injection of 150 μ g PAPP-A in Incomplete Freund's Adjuvant (ICA). One week later the mouse was boosted with an intravenous injection of 65 μ g PAPP-A and the animal sacrificed five days later.

Hybridomas were prepared by fusing splenocytes and
15 inguinal lymph node cells with X63 Ag 8.6.5.3. myeloma cells (Gaefee et al. 1977) and cultured in HAT medium. Once established, the selected hybridomas were cultured in aminopterin-free HT medium. Antibody production was monitored by enzyme immunoassay (EIA) and antibody
20 producing cells were cloned by limiting dilution and expanded. Hybridoma cells were injected intraperitoneally into BALB/c mice, primed with 2,6,10,14-tetramethyl pentadecane (pristene; Sigma, Sydney) to produce ascitic fluid.

25 Immunoassays:

Screening for antibody production was achieved by a crude EIA. Flat bottom Maxisorp Immunoplates (Nunc) were coated overnight (12-18h), at 4 C, with heparin-binding proteins (37.5 mg PAPP-A/well; See Isolation 1) extracted
30 from pooled pregnancy serum (38 weeks gestation: n 250). The plates were washed once with 50 mM sodium phosphate buffer, pH 7.4, containing 0.15M NaCl (PBS). Unreacted protein binding sites were blocked, by incubating for 90 min at room temperature (RT; 21-24 C), with PBS
35 containing 0.1% bovine serum albumin (BSA) and 0.2% H₂O₂. Plates were washed (X3) with PBS containing 0.1% BSA and 0.1% Tween-20 (T-PBS). Spent hybridoma culture media (250 μ l) was incubated for 2hr at room temperature after

which the plates were washed (X3) with T-PBS and incubated, for 1h at room temperature, with horseradish peroxidase conjugated rabbit anti-mouse IgG (200 μ l/well; Dakopatts, Sydney), diluted 1/5000 in T-PBS. The plates
5 were washed (X3) with T-PBS and incubated in the dark, for 15 min at RT, with substrate (200 μ l/well; O-phenylenediamine (OPD), 0.67 mg/ml), in 0.1M citric acid phosphate buffer, pH 5.0, containing 0.012% H₂O₂. The reaction was terminated by addition of 1M sulphuric acid
10 (100 μ l/well) and absorbance measured at 490 nm on a microplate reader (Biotek).

Positive hybridoma media were screened by PAPP-A RIA. The final incubation volume (200 μ l) consisted of hybridoma medium (100 μ l) and tracer (¹²⁵I-PAPP-A; 50 000
15 cpm/100 μ l). After overnight incubation at room temperature, pooled serum (50 μ l) obtained from normal human males was added to each tube and antibody bound tracer was precipitated by the second antibody-polyethylene glycol method (Sinosich *et al.*, 1982)⁹. The
20 supernatant was aspirated and precipitated radioactivity measured by a multichannel gamma-counter (NE, Sydney).

Murine immunoglobulins were typed by EIA using a panel of rabbit anti-mouse immunoglobulins (IgG₁, IgG_{2a}, IgG_{2b}, IgG₃, IgM, IgA, K light Chain, K Chain; Biorad
25 Laboratories, Sydney).

A rabbit polyclonal antibody was prepared by repeated intramuscular injections of pure PAPP-A (100 μ g). After a positive test bleed was obtained, the animal was boosted with 100 μ g PAPP-A administered
30 subcutaneously. Antibody responses were monitored by RIA on blood samples obtained from the marginal ear vein.

Biotinylation of Immunoglobulins:

Anti-PAPP-A IgG molecules were isolated from polyclonal and monoclonal antisera by affinity
35 chromatography on a 3 ml column of Protein-A Sepharose (Pharmacia, Sydney), as per manufacturer's instructions. Matrix bound proteins were desorbed with 0.2M glycine-HCl, pH 3.0, and dialysed (3 X 20 volumes)

against PBS. Approximately 1 mg of polyclonal IgG was recovered per ml of rabbit serum and 2.5 mg of IgG was recovered from 50 ml of culture supernate. Each preparation (1 mg/ml) was biotinylated with the long-chain derivative of N-hydroxysuccinimido-biotin (NHS-LC-biotin), as per manufacturer's instructions (Amersham, Sydney). After 1hr incubation at room temperature, unconjugated biotin was separated from the protein on a PD10 (Sephadex G-25M) column (Pharmacia) developed with PBS.

Development of PAPP-A Enzymeimmunoassay (EIA):

Micro-ELISA (96 well) plates were coated, overnight at 4 C, with polyclonal rabbit anti-PAPP-A IgG in PBS. After coating, the plates were washed with PBS and unreacted sites blocked (as detailed above). Assay reaction volume was 200 μ l with all samples/standards assayed in duplicate. Serum obtained from women (n>250) at advance stages of gestation (38 weeks) was pooled (LPS) and designated as 100 IU PAPP-A/L. The pooled LPS was serially diluted (1/16-1/8196) to prepare a standard curve, 12.2-6246 mIU/L).

Affinity Immuno-electrophoresis (AIE):

Pooled late pregnancy serum (38 weeks gestation; LPS) was incubated, overnight at room temperature, with media (control) or monoclonal antibody (5-62, 27-66, 25-1, 18-9) in a 1:1 ratio (v/v). An aliquot (10 μ l) was electrophoresed in the first dimension, in the absence and presence of heparin (+H; 20 units/ml gel), at 10 V/cm until the bromophenol blue marker migrated 3 cm. The gel was sliced and migrated in the second dimension into gel containing polyclonal rabbit anti-PAPP-A antibodies (Dakopatts) at 1/150 dilution. Second dimensional electrophoresis (at 2.5 V/cm) was performed overnight and the agarose gels processed for visualisation. Changes in migration distance (mm) were expressed as percentage of control value²⁷.

Immunohistochemistry (ICC):

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Term human placenta, obtained by caesarean section, was sliced and fixed in phosphate buffered formalin (4%), pH 7.4, for 24h at 4 C. The tissue was then washed in phosphate buffered saline, pH 7.4, for 24h and processed by standard histological techniques. Sections (5 μ m) were deparaffinised and processed for immunohistochemistry with polyclonal rabbit anti-chorionic gonadotrophin (Dakopatts), - PAPP-A (Sinosich et al. 1987) and monoclonal anti-PAPP-A (clone 25-1). Visualisation was achieved by enzyme bridge immunoperoxidase system using DAB as substrate. Negative controls included non-immune rabbit serum, mouse serum and culture media.

RESULTS

A total of five (5) anti-PAPP-A clones were obtained; 1) clone 25-1, isotyped as IgG₁ with K chain, 2) clone 27-66, isotyped as IgM with K chain, 3) clone 5-62, isotyped as IgG₁ with K chain, 4) clone 18-9, isotyped as IgG with K chain, and, 5) clone 41-1, which has yet to be characterised. By enzyme immunoassay, none of these antibodies reacted with immobilised heparin-Sepharose binding proteins extracted from normal male serum. The affinity constants (M^{-1}) for clones 25-1 and 5-62 were 1.26×10^{-9} and 2.65×10^{-9} , respectively.

A sandwich enzyme immunoassay was developed using immobilised rabbit polyclonal anti-PAPP-A immunoglobulins as capture phase and clone 25-1 immunoglobulins for PAPP-A quantification (Fig 3). This assay protocol is applicable to other PAPP-A monoclonal antibodies, but only clone 25-1 will be detailed.

Capture phase may include:

- 1) polyclonal PAPP-A antibody,
- 2) monoclonal PAPP-A antibody,
- 3) ligands, such as heparin, immobilised metal (zinc, copper, cobalt, nickel) cations and lectins (Con A, PHA) as detailed before,
- 4) specific, such as receptors, and non-specific,

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such as dyes, ligands for PAPP-A.

Coating concentrations of rabbit immunoglobulins ranged from 0.38-6.0 $\mu\text{g}/\text{well}$. At concentrations in excess of 1.5 $\mu\text{g}/\text{well}$, the increase in absorbance (at 490 nm) was insignificant. Thus, the EIA was developed on a coating concentration of 1.5 μg rabbit immunoglobulins per well. (See Figure 4).

The influence of time and temperature on PAPP-A capture by immobilised polyclonal anti-PAPP-A immunoglobulins is shown in Fig 5. Independent of PAPP-A doses (0-6250 mIU/L) and temperature (4°C; room temperature, 19-23°C; 37°C), binding equilibrium was attained by the capture antibody within 24h. At room temperature or 37°C, maximal PAPP-A capture was achieved after overnight incubation (15h) and these conditions (overnight incubation at room temperature) were selected for assay development.

Using biotinylated 25-1 antibody, at varied dilutions (1/5, 1/50, 1/500, 1/5000), binding equilibrium was achieved (at room temperature) by 4h. (Fig. 6) The time required to attain equilibrium was inversely proportional to antibody concentration, with maximal binding being achieved within 30 min in excess antibody (1/5 dilution). Using these conditions (4h at room temperature), Figure 7 shows typical PAPP-A dose-response curves at two dilutions (1/75, 1/150) of clone 25-1. For greater sensitivity the monoclonal antibody was diluted 1/75, but for routine assays not requiring enhanced sensitivity the antibody was diluted 1/150. Assay detection limit (n=20 replicates) was 21 mIU/L, with inter-assay precision of 6.2% and 10.5%. At PAPP-A doses of 100 and 1000 mIU/L and intra-assay precision was 9.2% (100 mIU/L) and 11.3% (1000 mIU/L), respectively. Total turnaround time for this assay was 24hrs.

Specificity was assessed in two ways. Firstly, by incubating each monoclonal PAPP-A antibody with immobilised heparin-Sepharose binding proteins, extracted from normal male serum. No significant reaction was

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detected. Secondly, at a fixed dose of PAPP-A (390.4 mIU/L) various monoclonal and polyclonal antisera (1/100 dilution) were tested for competition against biotinylated PAPP-A monoclonal 5-62, 25-1 (Table 1) and 18-9 (not shown). Positive reactions (>20% suppression in absorbance) were demonstrated with polyclonal anti-human, anti-monkey (rhesus) PAPP-A and rabbit anti-human placenta antisera (Table 2). Positive controls included the same monoclonal antibody as the test, but in the non-biotinylated form. All of the other antisera (PZP, a2M, SP1, complement factors 3 and 4, etc. Table 2) yielded negative results. Since the monoclonal anti-PAPP-A antibodies did not inhibit each other, except in a positive (self-self) format, these monoclonals recognise distinct epitopes.

Serially diluted late pregnancy serum, and extracts of placentae, delivered either by caesarean section or spontaneously per vagina, were analysed by RIA and EIA using clone 25-1 as primary antibody. Slopes of the logit-log regressed lines (Fig. 8 & 9) suggests parallelism between circulating and tissue PAPP-A. However, comparison of dose-response for extracts of human, chimpanzee and rhesus placentae showed marked species dependent responses. Clone 27-66 did not distinguish between three primate PAPP-A species (Fig.10). Clones 5-62 and 18-9 (not shown) demonstrated parallel lines only between human ($m = 1.82$) and chimpanzee ($m = 2.13$) PAPP-A (Fig. 11). In contrast, clone 25-1 reacted only with human ($m = 3.32$), but not with chimpanzee ($m = 0.36$) or rhesus ($m = 0.31$) PAPP-A (Fig 12).

After incubation with a variety of glycosidases immunoreactivity of radiolabelled (^{125}I -) PAPP-A to the two IgG clones (25-1, 5-62) was unchanged. Incubation with a-glucosidase and neuraminidase caused a marked increase in NSB, so that it was not possible to determine the true B_0 (%). By contrast, incubation of PAPP-A tracer with proteases (trypsin, elastase) markedly

reduced PAPP-A immunoreactivity. These findings suggest the monoclonal antibodies recognise protide epitopes, rather than carbohydrate moieties.

By two dimensional immunoelectrophoresis, PAPP-A had a 2-electrophoretic mobility (39 mm). In the presence of IgG (5-62, 25-1, 18-9) and IgM (27-66) clones, PAPP-A mobility was reduced by 25.6% and 43.6%, respectively. In the presence of heparin (20 units/ml gel), PAPP-A migration was increased by 33.3%. As none of the four clones inhibited the heparin-PAPP-A interaction, these epitopes are not associated with the heparin binding site of PAPP-A.

Immunohistochemical localisation of PAPP-A (clone 25-1) in human term placenta was limited to the syncytial layer. No PAPP-A immunoreactivity was detected in foetal or maternal blood vessels or villous stroma.

The monoclonal antibody is typically used for diagnostic or therapeutic purposes in unlabelled and/or labelled form. Typical labels include fluorescent, biotin, radioisotopic, chromogenic, chemiluminescent, electron dense, magnetic labels and enzyme labels, as examples. Labelling may also be achieved by means of a second antibody against the primary antibody with the second antibody being detectably labelled.

PAPP-A measurement in female (ovarian follicular fluid, cervical mucus) and male (seminal plasma, prostatic fluid) reproductive tract secretions can be of use to assess folliculogenesis, granulosa cell status, ovulation and pathology of the male accessory glands, such as prostate gland. Each fluid should be centrifuged to remove cells and the supernate stored at -20°C, until required for analysis. Cervical mucus should be extracted with 500 ul of assay buffer, centrifuged to remove cells and cellular debris, and supernate stored at -20°C until required for analysis.

Blood, obtained by routine venepuncture, can be collected into anticoagulant free tubes or into anticoagulant (heparin, EDTA, citrate, as examples)

containing tubes. In each case the sample should be centrifuged to separate the serum/plasma phase from blood cells. The supernate fluid is aspirated and stored at -20°C until required for analysis.

5 In the case where trophoblast cells are to be isolated, anticoagulated (EDTA, citrate, heparin) maternal blood or vaginal fluid is washed with physiological buffers or culture media to remove the plasma. The sedimented cells are resuspended in buffer
10 or culture media and incubated with one or more PAPP-A polyclonal or monoclonal antibodies (or their derivatives). These PAPP-A antibodies may be free or immobilised, for example on magnetised particles. The free PAPP-A antibodies can be removed with a capture
15 phase, such as Protein-A Sepharose, secondary antibody immobilised on magnetic and non-magnetised particles. The isolated trophoblast cells can be used for fetal karyotype, fluorescent in situ hybridisation (FISH); molecular biology, cell culture and immunocytochemical
20 studies.

Tissue, for cell culture and immunohistochemical analyses, can include any desired normal and pathological tissue. Cell isolation and tissue processing was achieved by standard protocols as detailed above.

25 Immunodetection of PAPP-A, in maternal blood, can be used to assess pregnancy wellbeing and as part of a prenatal biochemical screen to assess fetal development. In all adverse clinical situations, such as extra-uterine pregnancies, fetal death in utero, anembryonic pregnancy,
30 incomplete abortion, spontaneous miscarriage, fetal malformation and/or aneuploidy (chromosomal abnormality), as examples, PAPP-A levels are depressed in maternal circulation. Although, in most of the pregnancy disorders, hCG (or free β -subunit) expression correlates
35 with PAPP-A levels. In Trisomy 21 (Down Syndrome) hCG (or free β -subunit) expression is increased. Therefore, development and application of an algorithm which combines PAPP-A and hCG (or free β -subunit) measurements

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will permit discrimination of Trisomy 21 (Down Syndrome) from other pregnancy disorders. With a sensitive immunoassay, PAPP-A measurement can be established as an integral part of antenatal care in the first trimester of pregnancy.

In oncological situations, PAPP-A is expressed by tumor granulosa cells and gestational trophoblastic diseases (GTD). As the GTD becomes more aggressive and metastatic, PAPP-A expression is downregulated. By contrast, hCG and free β -subunit expression is upregulated. Again, application of a PAPP-A - hCG (or free β -subunit) algorithm enables discrimination between malignant and benign GTD.

INDUSTRIAL APPLICABILITY

In addition to the industrial applicability already indicated, it is envisaged that PAPP-A and PAPP-A variants can be used as a medicament to treat pregnant women exhibiting low levels of PAPP-A.

The amount of PAPP-A or variant required to produce a single dosage form will vary depending upon the condition to be treated, patient to be treated and the particular mode of administration. The specific dose level for any particular patient will depend upon a variety of factors including the activity of the molecule employed, the age, body weight, general health and diet of the patient, time of administration, route of administration, rate of excretion, drug combination and the severity of the condition undergoing treatment.

TABLE 1LIST OF ANTIBODIES CHECKED FOR CROSSREACTIVITY AGAINST THE PAPP-A MONOCLONALSDAKO ANTIBODIES

5 alphafetoprotein, albumin, al-antichymotrypsin, antithrombin 111, al-antitrypsin, C reactive protein, C1 esterase inhibitor, C3C complement, C3C(unspecific), C3d complement, C4 complement, cholinesterase, fibrinogen, fibronectin, human chorionic gonadotropin, human placental lactogen, inter-a-trypsin inhibitor, B-lipoprotein, a2-macroglobulin orosomucoid, PAPP-A, a2-plasmin inhibitor, plasminogen, pregnancy zone protein, schwangerschaftsprotein 1, placenta and B2-microglobulins.

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PROTEIN A-ISOLATED RABBIT SERUM

Prenancy protein 12, pregnancy-associated plasma protein B, and chimp PAPP-A.

RABBIT SERUM

15 Human serum PAPP-A 1&2, chimp PAPP-A, rhesus PAPP-A, heparin-bound macaque LPS, macaque male serum, heparin-bound human male serum, heparin bound guinea pig placental homogenate, guinea pig placental proteins 1&2, seminal plasma, pregnancy protein 10, pregnancy protein 14, pregnancy protein 5, and inhibin.

MONOCLONAL ANTIBODY SUPERNATANTS

20 25-1 IgG, 25-35 IgG,¹ 27-66 IgM, 27-105 IgM,¹ 5-62 IgG, 5-12 IgG,¹ 5-39 IgG,¹ 5-17 IgG,¹ 11-1-30 IgM,²

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TABLE 2

SPECIFICITY	% REDUCTION IN O.D.	
	25-1	5-62
Dako anti-PAPP-A	-78.4%	-91.9%
Dako anti-Placenta	-47.8%	-56.2%
Rabbit anti-Human serum PAPP-A 1	-71.3%	-96.1%
Rabbit anti-Human serum PAPP-A 2	-72.6%	-90.5%
Rabbit anti-Rhesus PAPP-A	-38.8%	-86.3%
Protein-A isolated anti-Chimp PAPP-A	-60.4%	-29.9%
25-1 Monoclonal antibody supernatant	-50.8%	-
25-35 Monoclonal antibody supernatant	-46.6%	-
5-62 Monoclonal antibody supernatant	-	-61.7%
5-23 Monoclonal antibody supernatant	-	-60.8%
5-39 Monoclonal antibody supernatant	-	-62.6%
5-17 Monoclonal antibody supernatant	-	-59.5%

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Claims:

1. The protein PAPP-A, substantially free of other human proteins, which is a homotetramer having a molecular weight of approximately 820 kiloDaltons and an isoelectric point of approximately 4.5, each monomeric subunit of the homotetramer having a molecular weight of approximately 200 kiloDaltons and an N-terminal amino-acid sequence of Glu-Ala-Arg-Gly-Ala-Pro-Glu-Glu-Pro-Ser-Pro-Pro-Ser.
 2. A variant of the protein according to claim 1.
 3. A polynucleotide encoding PAPP-A according to claim 1 or a PAPP-A variant according to claim 2.
 4. A method for purifying PAPP-A from a sample comprising the steps of:
 - (i) applying the sample to a positive affinity chromatography on Heparin-Sepharose; and
 - (ii) size fractionating the fractions obtained from step (i) on a gel filtration column; and
 - (iii) applying the fractions obtained from step (ii) to an anion-exchange column; and
 - (iv) applying the fractions obtained from step (iii) to a matrix with immobilized antibodies.
- Steps (i), (ii) and (iii) can performed in any sequence. The above is merely one protocol.
5. The method according to claim 4 wherein the steps (i), (ii) and (iii) can be performed in any order.
 6. The method according to claim 4 further including applying the fractions obtained from the matrix to a Heparin-Sepharose column.
 7. A monoclonal antibody which recognises PAPP-A.
 8. A monoclonal antibody which recognises PAPP-A but does not significantly cross-react with immobilised heparin-Sepharose binding proteins extracted from normal male serum.

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9. A monoclonal antibody specific for PAPP-A.
10. A monoclonal antibody according to any one of claims 7, 8 or 9 which is an IgG or IgM antibody.
11. A monoclonal antibody according to any one
5 of claims 7, 8 or 9 which has kappa light chains.
12. A monoclonal antibody produced by clones 25-1, 27-66, 5-62, 18-9 or 41-1, as herein before defined, or an antibody, or fragments thereof, which recognises the same epitope as an
10 a n t i b o d y produced by one of the clones.
13. An assay for PAPP-A comprising using a monoclonal antibody according to any one of claims 7 to 12 to detect PAPP-A.
14. An assay according to claim 13 which is an
15 ELISA assay, immunoradiometric assay or any other assay format.
15. An assay according to claim 13 comprising a specific or a non-specific ligand as capture phase.
16. An assay according to claim 13 wherein the
20 monoclonal antibody may or may not be directly tagged.
17. A method of detecting PAPP-A which comprises the steps of:
- (a) contacting the sample with a PAPP-A capture
phase which consists of one or more of the
25 following:
- (i) monoclonal PAPP-A antibody;
 - (ii) polyclonal PAPP-A antibody;
 - (iii) immobilised heparin;
 - (iv) immobilised divalent metal cations
30 (Cu⁺⁺, Zn⁺⁺, Co⁺⁺, Ni⁺⁺),
 - (v) immobilized lectins (Concanavalin A, Helix pomatia, Lens culinaris, Limulus polyphemus, phytohaemagglutinin, Ricinus communis, Wheat germ, or others with
35 equivalent specificities), and
 - (vi) other specific (receptors) and non-specific (dyes) ligands.
- (b) contacting the immobilised or captured PAPP-A

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sample with a monoclonal antibody according to anyone of claims 6 to 11 labelled with a detectable marker;

(c) incubating the sample and the labelled antibody to permit the labelled antibody to bind to any PAPP-A in the sample and;

(d) detecting the labelled protein.

18. A kit for the detection of PAPP-A in a sample,, said kit comprising at least one monoclonal antibody according to any one of claims 7 to 12 together with a positive control or a negative control.

19. Use of PAPP-A as a medicament together with a pharmaceutically acceptable excipient to treat a pregnant patient exhibiting the absence of or low levels of PAPP-A.

20. A medicament comprising an effective amount of PAPP-A together with a pharmaceutically acceptable carrier, diluent or excipient.

21. Use of PAPP-A as a contraceptive agent.

22. Use of PAPP-A antibodies to immunoneutralise unwanted conceptions abnormal pregnancies and trophoblastic tumors.

23. Use of PAPP-A antibodies for the isolation of trophoblast cells, from maternal circulation or reproductive tract, for prenatal fetal karyotyping.

24. Use of PAPP-A antibodies for the diagnosis and of oncological disorders and compromised pregnancies.

25. Use of PAPP-A antibodies for prenatal screening for Down Syndrome and diagnosis of malignant trophoblastic disease.

26. An epitope of PAPP-A, being an epitope recognised by at least one of the monoclonal antibodies of claims 7 to 12.

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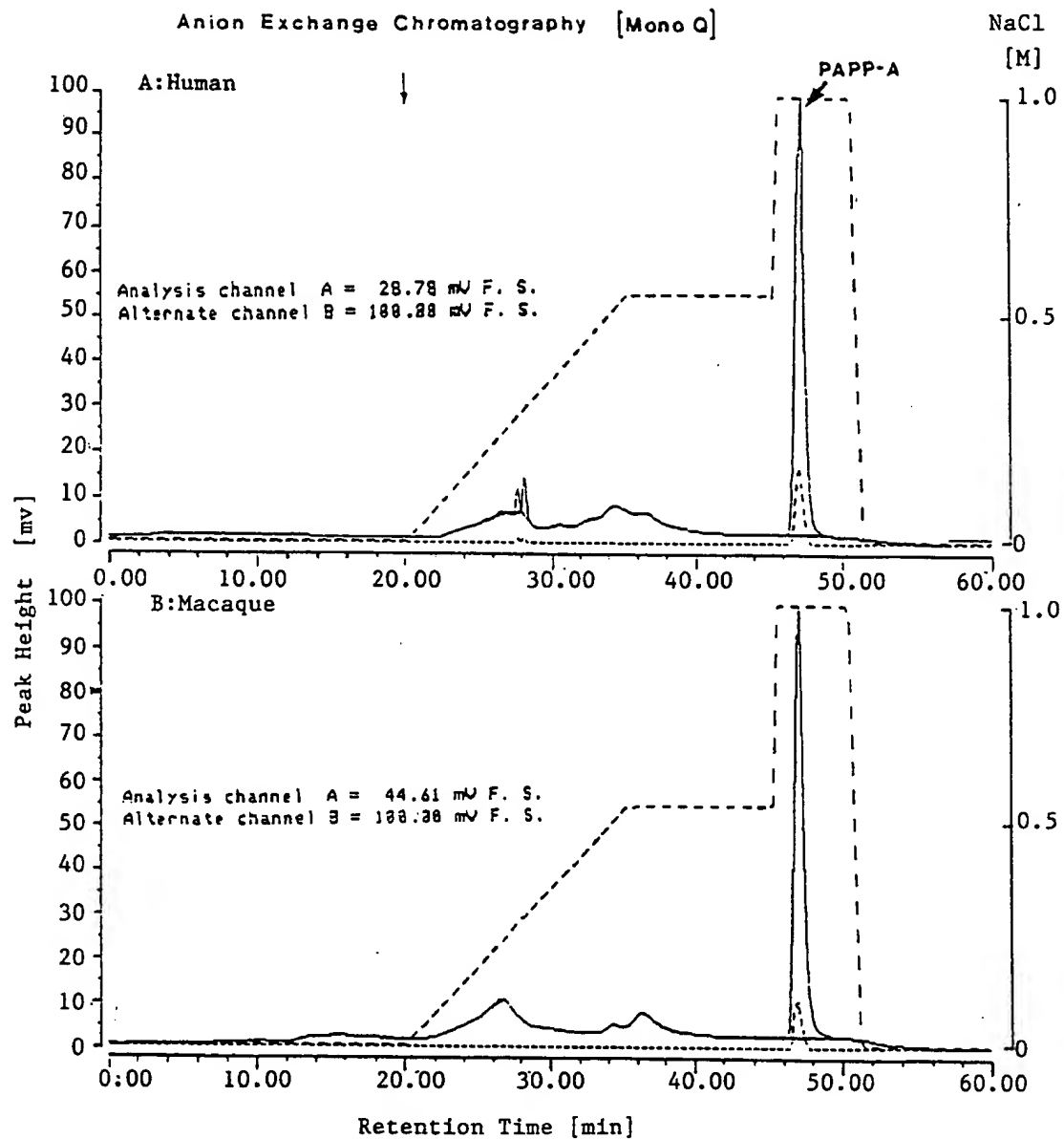
FIGURE 1

N-terminal amino acid sequence of human PAPP-A.

Glu-Ala-Arg-Gly-Ala-Pro-Glu-Glu-Pro-Ser-Pro-Pro-Ser

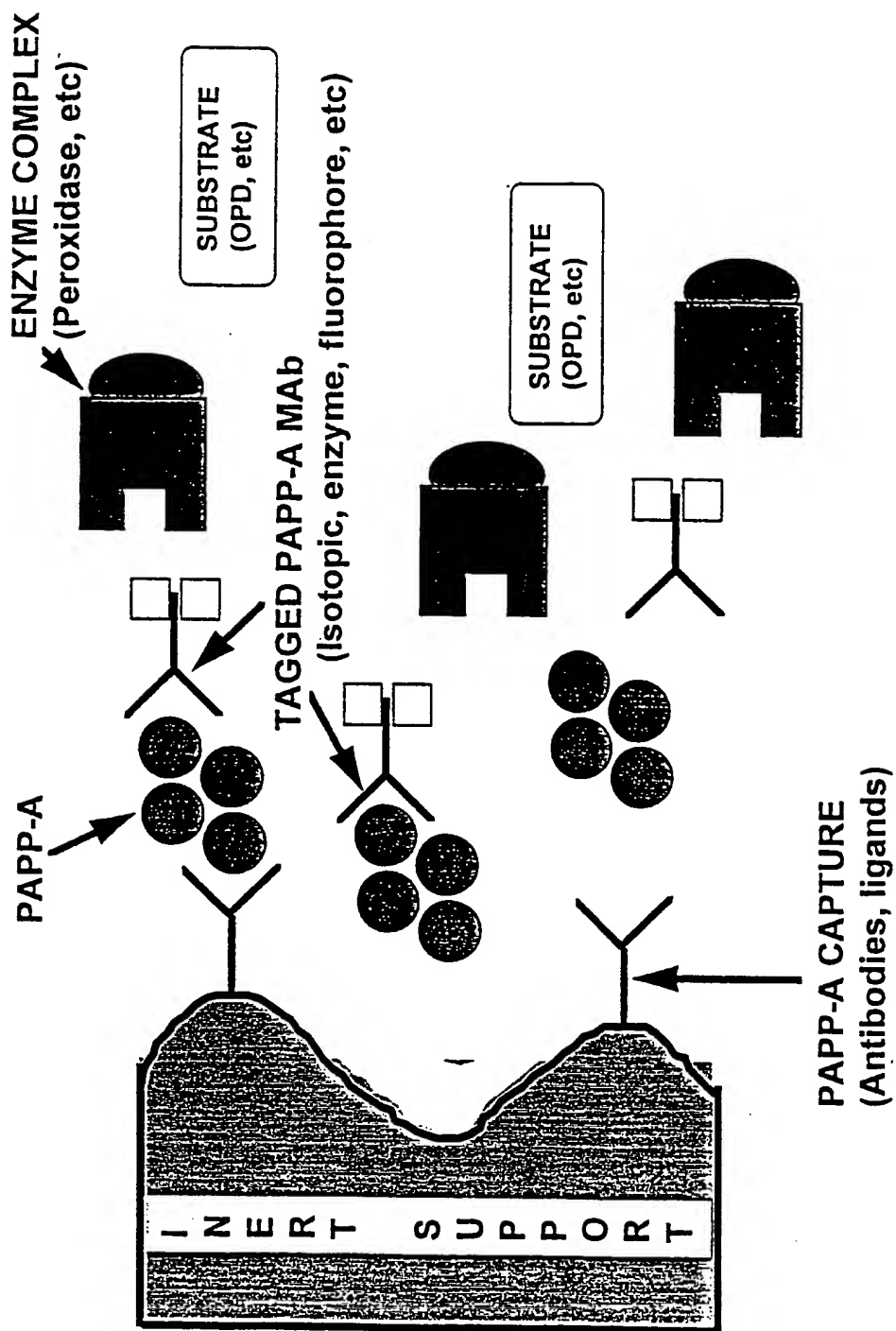
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FIGURE 2



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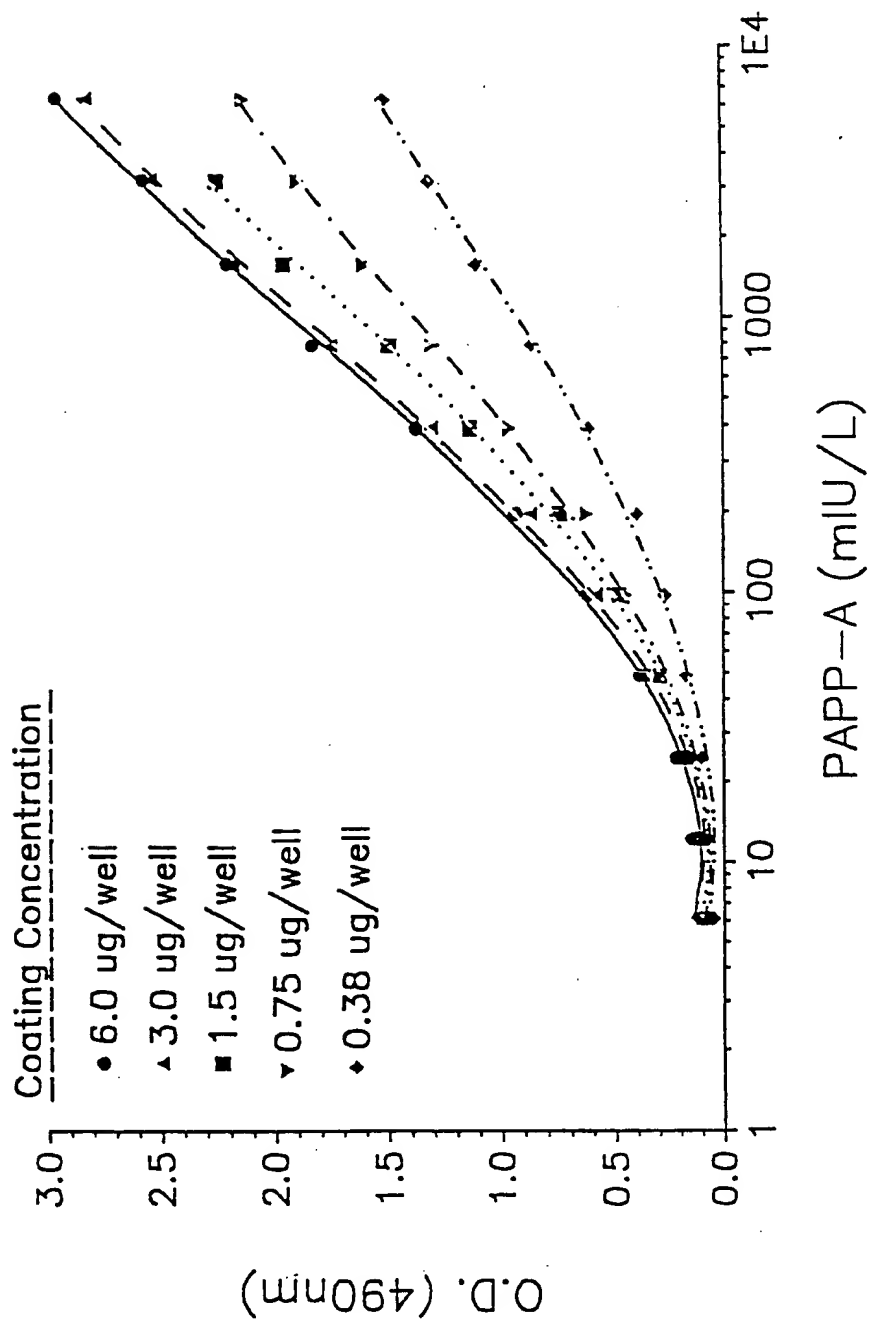
FIGURE 3

*Figure 3 Sandwich PAPP-A assay formats*

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FIGURE 4

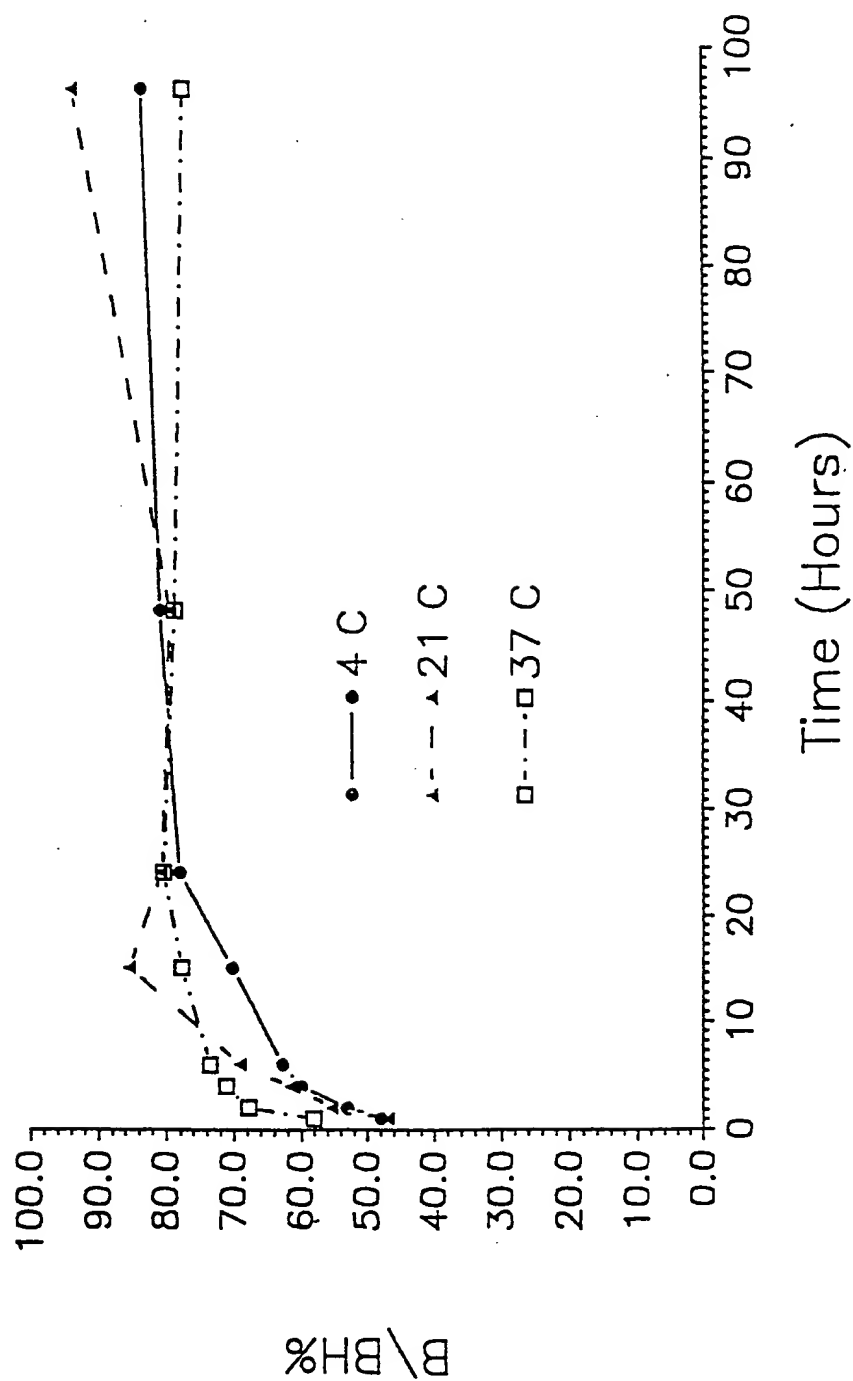
Standard Curves for PAPP-A



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FIGURE 5

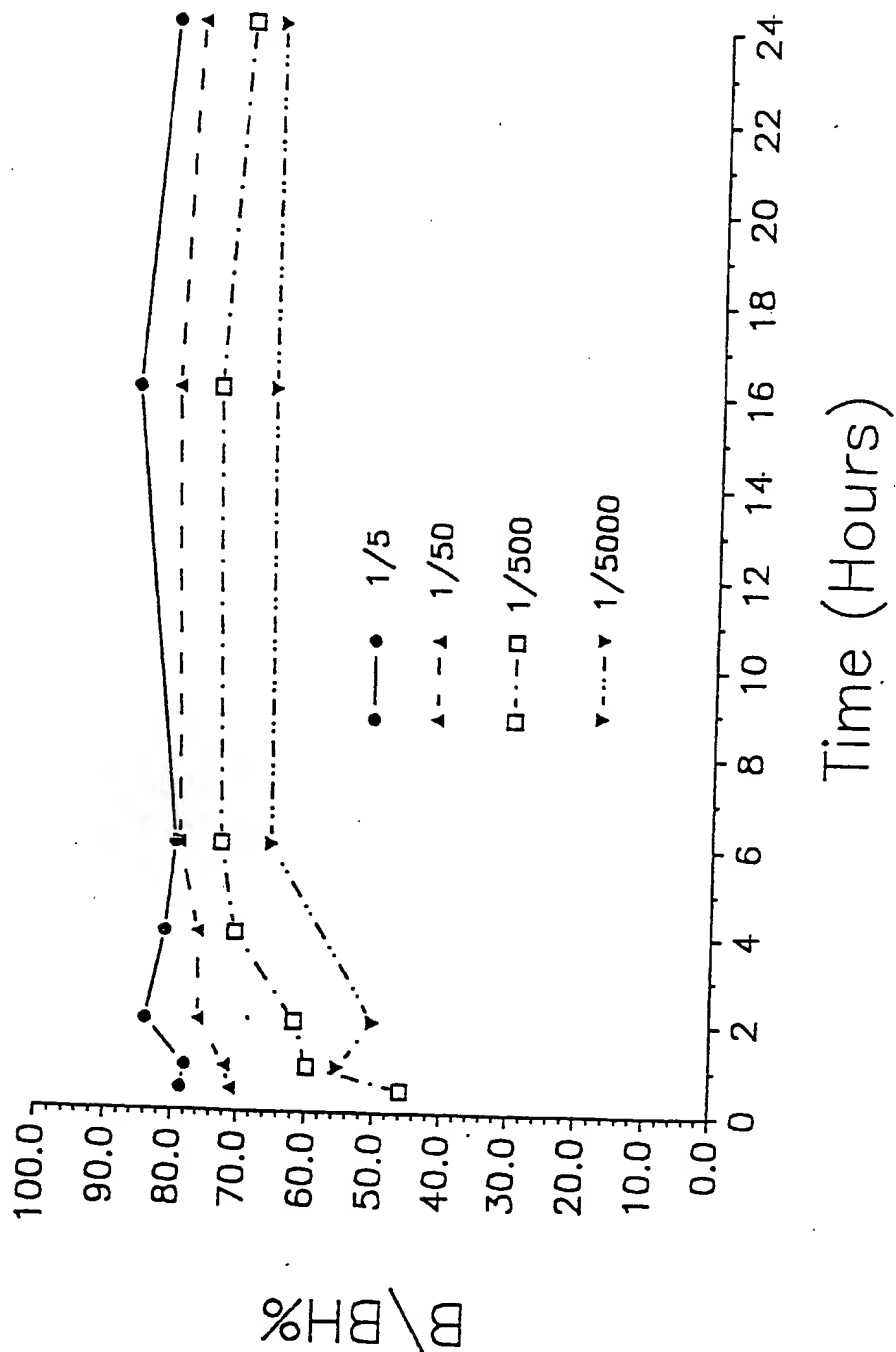
Influence of Temperature on Binding Equilibrium of Immobilised Immunoglobulin



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FIGURE 6

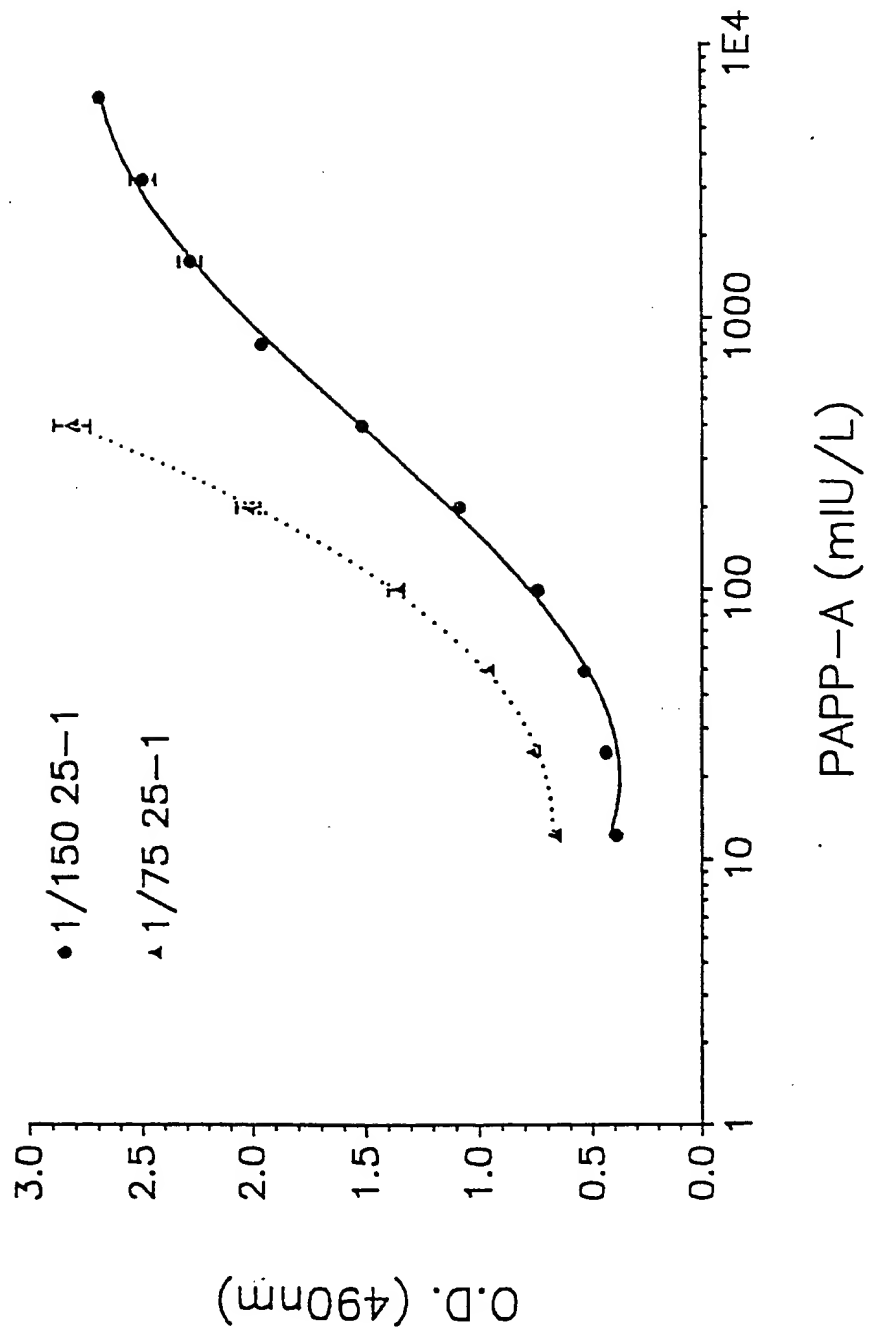
Binding Equilibrium of PAPP-A Monoclonal Antibody (25-1)



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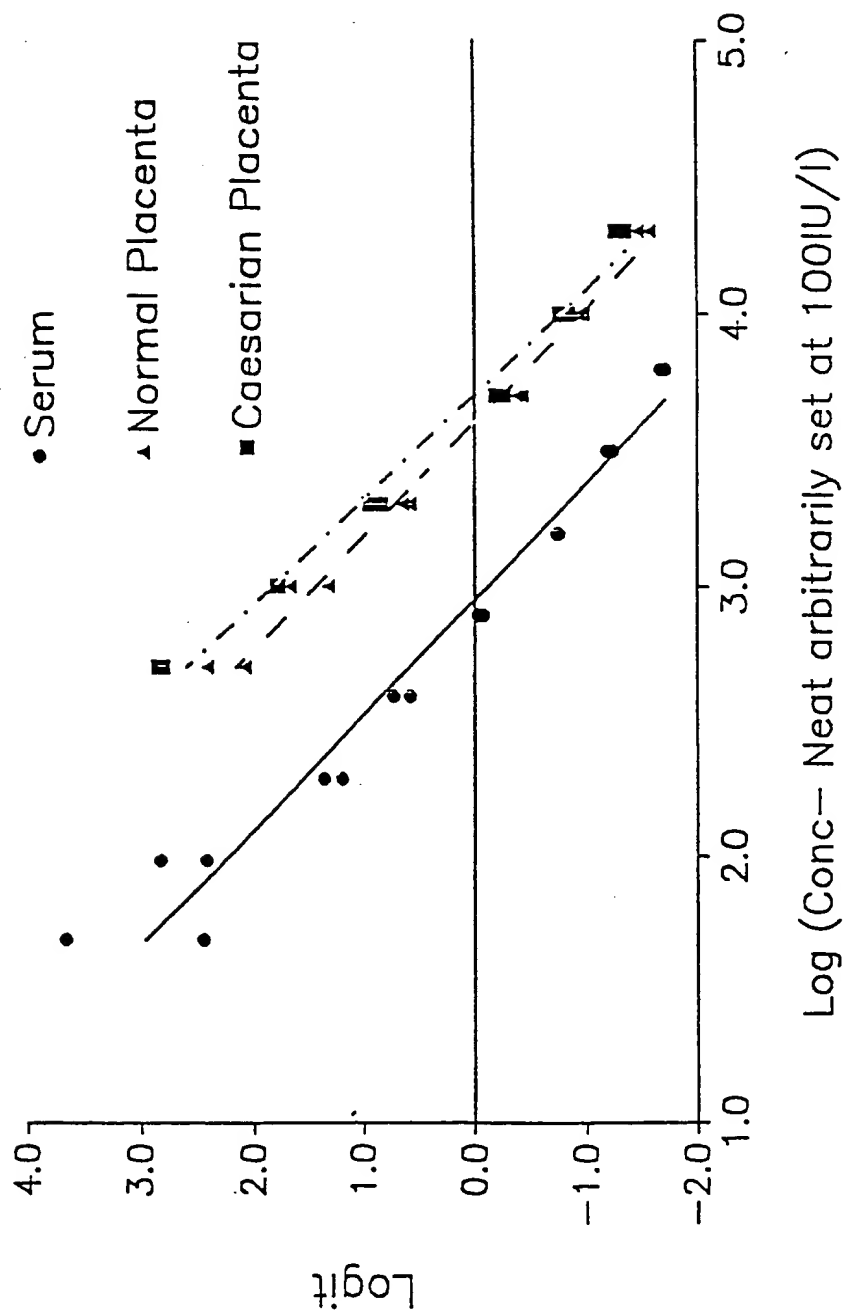
FIGURE 7

Standard curves for PAPP-A



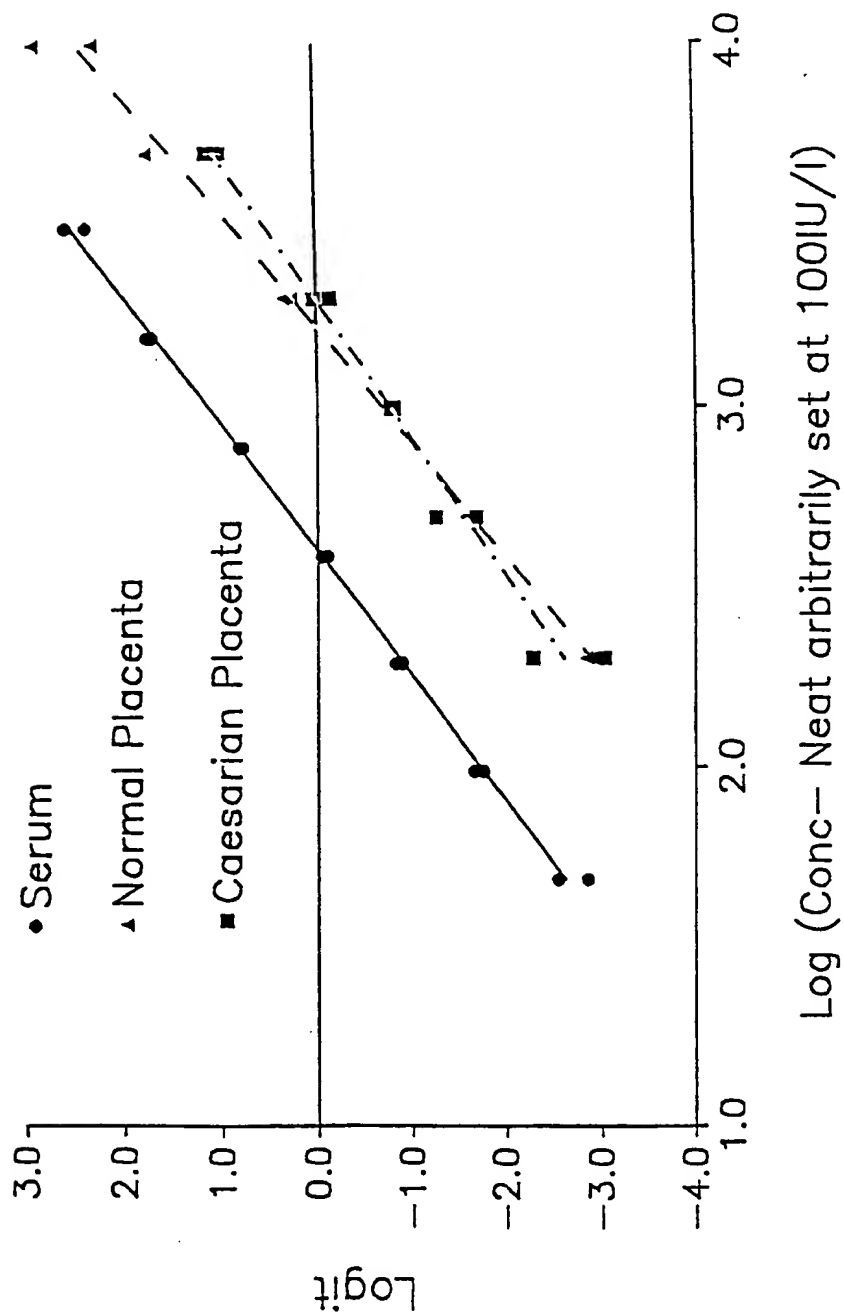
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FIGURE 8

Logit Log Graphs for Human Serum and Normal and
Caesarian Placental Homogenates assayed by RIA



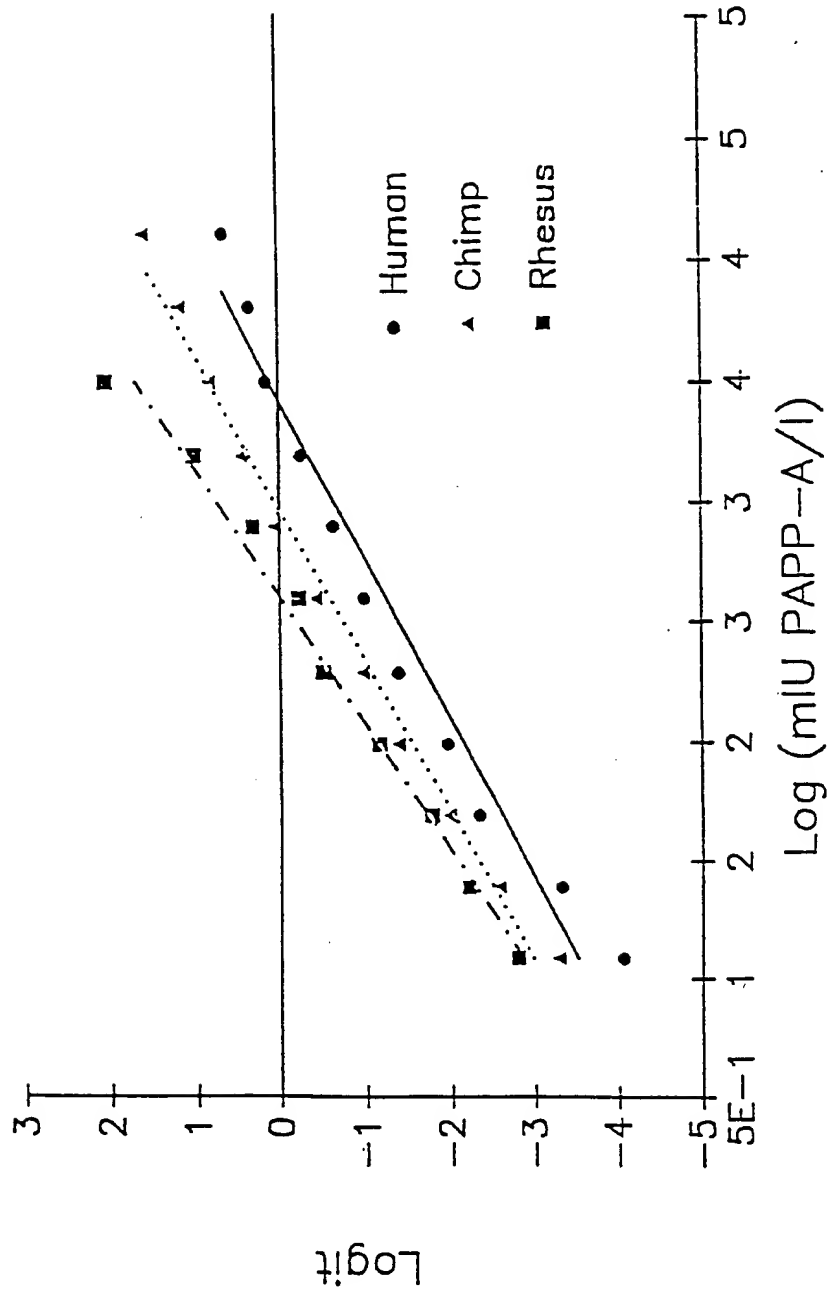
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FIGURE 9

Logit Log Graphs for Human Serum and Normal and
Caesarian Placental Homogenates assayed by EIA with 25--1



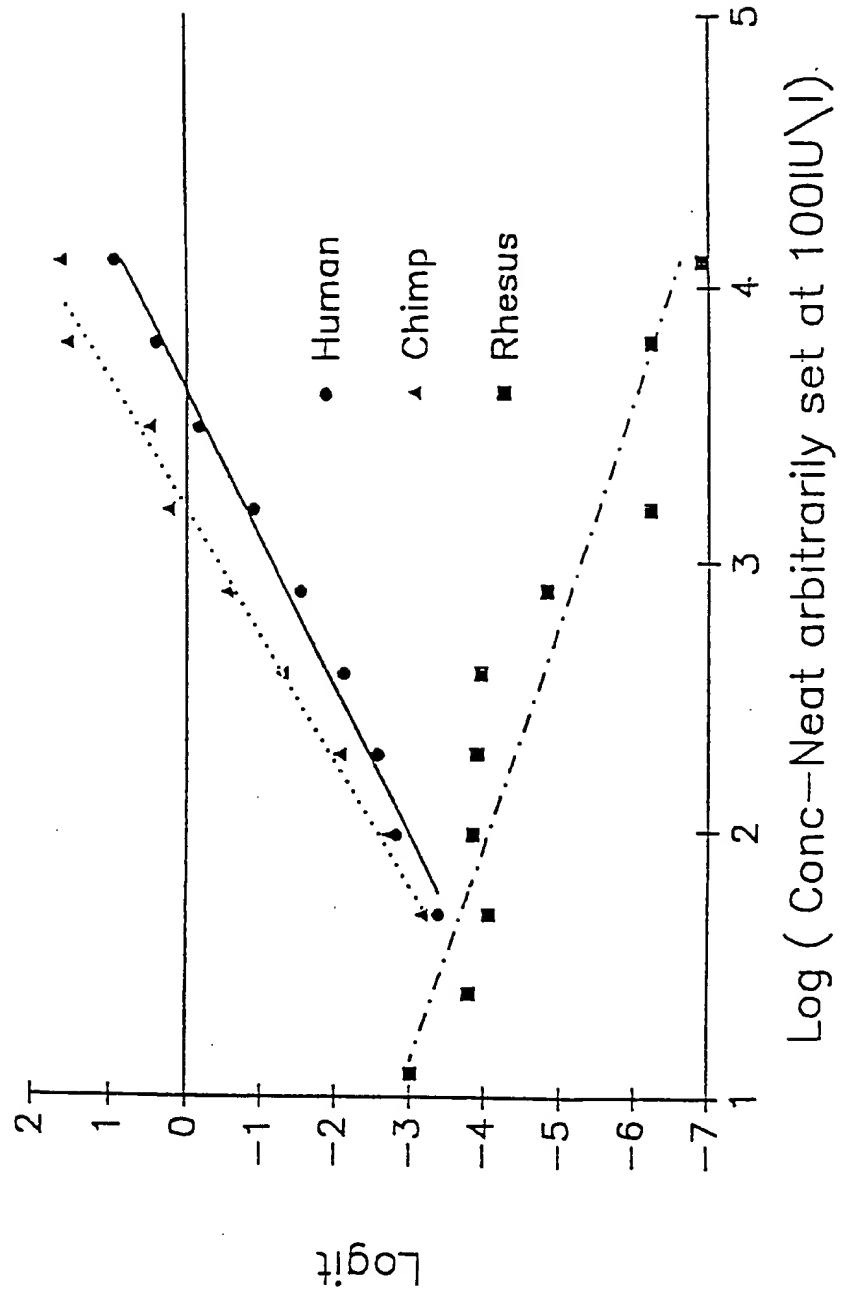
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FIGURE 10

Logit-Log Graphs for Human and Chimp Placental
Homogenates assayed by EIA with 27-66



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FIGURE 11

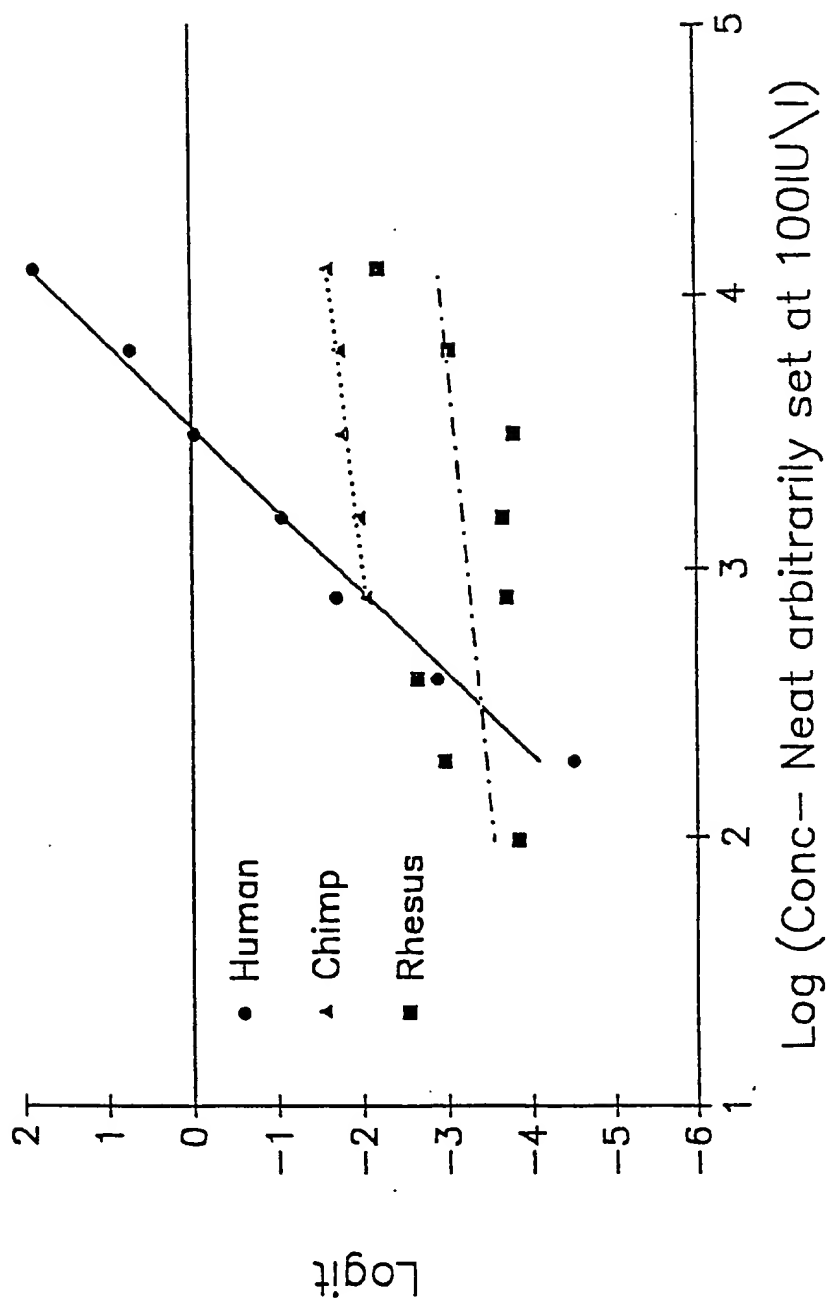
Logit-Log Graphs for Human and Chimp and Rhesus
Placental Homogenates assayed by EIA with 5-62



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FIGURE 12


Logit-Log Graphs for Human and Chimp Placental
Homogenates assayed by EIA with 25-1



INTERNATIONAL SEARCH REPORT

International application No.

PCT/AU 94/00139

A. CLASSIFICATION OF SUBJECT MATTER Int. Cl. ⁵ C07K 15/06, 15/14, 15/12; C12P 21/08; G01N 33/577; A61K 27/02, 39/395 According to International Patent Classification (IPC) or to both national classification and IPC				
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) IPC: DERWENT AS BELOW Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched AU: IPC C07K 7/10, 13/00, 15/06, 15/14 Electronic data base consulted during the international search (name of data base, and where practicable, search terms used) CHEMICAL ABSTRACTS (CAS): PAPP or PREGNAN: and ASSOCI: and PLASMA: and PROTEIN: DERWENT: PAPP or PREGNAN: and PLASMA: and PROTEIN: STN REGISTRY FILE: "EARGAPEEPSPPS" - SEQUENCE SEARCH				
C. DOCUMENTS CONSIDERED TO BE RELEVANT				
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to Claim No.		
X	AU-B-25177/88 (620351) (ADEZA BIOMEDICAL CORP) 20 February 1992 (20.02.92) see whole document	7, 9-11, 13-18, 24		
X Y	GYNECOL OBSTET INVEST Volume 32 No. 2 pages 72-77 (1991) M J SINOSICH et al. "PREGNANCY-ASSOCIATED ..."	1, 7, 9-11, 13-18 4-6		
X	BIOLOGY OF REPRODUCTION Volume 44 No. 1 pages 201-206 (1991) N CHEGINI et al. "THE PRESENCE OF PREGNANCY-ASSOCIATED PLASMA ..."	1, 7, 9-11, 13-18		
<div style="display: flex; justify-content: space-between;"> <div style="width: 45%;"> <input checked="" type="checkbox"/> Further documents are listed in the continuation of Box C. </div> <div style="width: 45%;"> <input checked="" type="checkbox"/> See patent family annex. </div> </div>				
<table style="width: 100%; border: none;"> <tr> <td style="width: 50%; vertical-align: top;"> <p>* Special categories of cited documents :</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> </td> <td style="width: 50%; vertical-align: top;"> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</p> <p>"&" document member of the same patent family</p> </td> </tr> </table>			<p>* Special categories of cited documents :</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p>	<p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</p> <p>"&" document member of the same patent family</p>
<p>* Special categories of cited documents :</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p>	<p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</p> <p>"&" document member of the same patent family</p>			
Date of the actual completion of the international search 6 July 1994 (06.07.94)		Date of mailing of the international search report 14 JUL 94 (14.07.94)		
Name and mailing address of the ISA/AU AUSTRALIAN INDUSTRIAL PROPERTY ORGANISATION PO BOX 200 WODEN ACT 2606 AUSTRALIA Facsimile No. 06 2853929		Authorized officer  T. SUMMERS Telephone No. (06) 2832291		

INTERNATIONAL SEARCH REPORT

International application No.

PCT/AU 94/00139

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate of the relevant passages	Relevant to Claim No.
X	ARCH GYNECOL OBSTET 247(2) pages 53-62 (1990) M J SINOSICH et al. "A-baboon model for pregnancy-associated ... "	1, 7, 9-11, 13-18
X Y	BIOCHEMISTRY INTERNATIONAL 20(3) pages 579-89 (1990) M J SINOSICH et al. "Characterisation of Pregnancy ... "	1, 7, 9-11, 13-18 4-6
X Y	ELECTROPHORESIS, 11(1), pages 70-8 (1990) M J SINOSICH "Molecular characterization of Pregnancy-associated ... "	1, 2
X Y	PLACENTA, 10(6) pages 569-78 (1989) M J SINOSICH et al. "RU-486 Induced Suppression of Placental Neutrophil ... "	1, 7, 9-11, 13-18 4-6
X Y	Br J Obstet Gynaecol, 96(7), pages 870-5 (1989) P Bischof et al. "Immunological heterogeneity ... "	1, 7, 9-11, 13-18 4-6
X	J Immunol Methods, 95(1), pages 129-33 (1986) E A Mowles et al. "A two site immunoradiometric ... "	7, 9-11, 13-18
X Y	Arch Gynecol, 236, (2), pages 83-91 (1984) L G Pinto Furtado et al. "The development and validation of ... "	1, 7, 9-11, 13-18 4-6
X Y	Anal Biochem, 131(1), pages 18-24 (1983) M W Davey et al. "Interaction between Heparin and Human ... "	1, 7, 9-11, 13-18 4-6
A	J Clin Invest, 54, (3), pages 576-82 (1974) Lin Tsue-Ming et al. "Measurement of pregnancy ... "	
P,X P,Y	Biochem, 33, (6), pages 1592-8 (1994) J Kristensen et al. "Amine Acid Sequence of Human Pregnancy ... "	1, 2 7, 9-11, 13-18

Information on patient family members

PCT/AU 94/00139

Patent Document Cited in Search Report		Patent Family Member	
AU-B 25177/88	EP-A2 316919 US-A 5223440 CA-AA 2098180 US-A 5281522	US-A 5096830 JP-A2 1195848 US-A 5236846	EP-A1 563165 US-A 5185270 WO-A1 9210585

END OF ANNEX